(19) World Intellectual Property Organization

International Bureau





(43) International Publication Date 28 September 2006 (28.09.2006)

PCT

(10) International Publication Number WO 2006/101706 A1

(51) International Patent Classification:

A61K 31/726 (2006.01) **A23L 2/52** (2006.01) **A61K 31/7008** (2006.01) **A23L 1/05** (2006.01)

(21) International Application Number:

PCT/US2006/007921

(22) International Filing Date: 6 March 2006 (06.03.2006)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:

60/662,705 17 March 2005 (17.03.2005) US

(71) Applicant (for all designated States except US): CARGILL, INCORPORATED [—/US]; 15407 McGinty Road West, Wayzata, Minnesota 55391 (US).

- (72) Inventors; and
- (75) Inventors/Applicants (for US only): JOHNSON, Janice [US/US]; 8684 Bayard Court, Eden Prairie, MN 55347 (US). HUNSICKER, John [US/US]; 1930 Girard Avenue South, Minneapolis, Minnesota 55403 (US). CROSBY, Laura [US/US]; 6301 Rowland Road, Eden Prairie, Minnesota 55344 (US). NELSON, Karla, Jean [US/US]; 4838 Aldrich Avenue South, Minneapolis, Minnesota 55419 (US).
- (74) Agents: LEVINE, Edward, L. et al.; CARGILL, IN-CORPORATED, 15407 McGinty Road West, Wayzata, MN 55391 (US).

- (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: HEALTH-RELATED LIQUID COMPOSITIONS

(57) Abstract: There are disclosed liquid compositions comprising water, a fiber such as inulin, and a glycosaminoglycan that is selected from chondroitin sulfate, glucosamine, or a mixture thereof. The liquid compositions may further comprise at least one or more sweeteners, a source of calcium, at least one or more flavor additives, trehalose, at least one or more acids, sodium citrate, monopotassium phosphate, fruit juice, fruit juice concentrate, or the like, and any combination thereof. The liquid compositions may be suitable for use in health-related applications, among other applications.



HEALTH-RELATED LIQUID COMPOSITIONS

FIELD OF DISCLOSURE

The present disclosure relates to health-related liquid compositions that are expected to provide, for example, joint health benefits.

BACKGROUND

Health-related liquid compositions are known. However, there is a continuing need to provide health-related liquid compositions that offer other benefits.

10

15

20

25

30

5

SUMMARY OF THE DISCLOSURE

This disclosure relates to liquid compositions comprising water, a soluble fiber, such as preferably inulin, and chondroitin sulfate or glucosamine, or both. Other ingredients such as a source of calcium, a flavor, a sweetener, citric acid, malic acid, sodium citrate dihydrate, monopotassium phosphate, trehalose, fruit juice, fruit juice concentrate, and the like, may be incorporated into the liquid compositions.

DETAILED DESCRIPTION OF THE DISCLOSURE

This disclosure relates to liquid compositions comprising water, a soluble fiber such as preferably inulin and chondroitin sulfate or glucosamine or both. Other ingredients such as a source of calcium, a flavor, a sweetener, citric acid, malic acid, sodium citrate dihydrate, monopotassium phosphate, trehalose, fruit juice, fruit juice concentrate, and the like, may be incorporated into the liquid compositions.

In one embodiment, the disclosure relates to a liquid composition comprising water, a soluble fiber such as inulin, and chondroitin sulfate. Other ingredients such as sweeteners, sources of calcium such as calcium lactate, trehalose, flavors, citric acid, malic acid, sodium citrate, monopotassium phosphate, fruit juice, fruit juice concentrate, and the like, may be incorporated into the liquid composition.

In another embodiment, the disclosure relates to a liquid composition comprising water, a soluble fiber such as inulin, and glucosamine. Other ingredients such as sweeteners, sources of calcium such as calcium lactate, trehalose, flavors, citric acid, malic

acid, sodium citrate, monopotassium phosphate, fruit juice, fruit juice concentrate, and the like, may be incorporated into the liquid compositions.

In another embodiment, the disclosure relates to a liquid composition comprising water, a soluble fiber such as inulin, chondroitin sulfate, and glucosamine. Other ingredients such as sweeteners, sources of calcium such as calcium lactate, trehalose, flavors, citric acid, malic acid, sodium citrate, monopotassium phosphate, fruit juice, fruit juice concentrate, and the like, may be incorporated into the liquid composition.

5

10

15

20

25

30

In another embodiment, the disclosure relates to a liquid composition comprising about 94.7% water, about 2.0% erythritol, about 1.4% inulin, about 0.78% calcium lactate, about 0.5% trehalose, about 0.08% natural lemon, about 0.08% natural lime, about 0.05% citric acid, about 0.05% malic acid, about 0.04% sodium citrate, dihydrate, about 0.02% acesulfame potassium artificial sweetener, about 1% solution, about 0.01% monopotassium phosphate, and about 0.17% chondroitin sulfate.

In another embodiment, the disclosure relates to a liquid composition comprising about 94.7% water, about 2.0% erythritol, about 1.4% inulin, about 0.78% calcium lactate, about 0.50% trehalose, about 0.13% peach flavor, about 0.05% citric acid, about 0.05% malic acid, about 0.04% sodium citrate, dihydrate about 0.02% acesulfame potassium artificial sweetener, 1% solution, about 0.01% monopotassium phosphate, and about 0.17% chondroitin sulfate.

In the embodiments of the disclosure, the following is applicable.

Any source of soluble fiber may be utilized in producing the liquid compositions. Conventional sources of soluble fibers are gums, hydrocolloids, mucilages, hemicellulose, pectins, and the like, and mixtures thereof. Suitable sources of soluble fiber include, but are not limited to, gum arabic, locust bean gum, polydextrose, inulin, arabinogalactose, hydrolyzed guar gum, xanthan, pectin, alginate, carrageenan, beta-glucans, tragacanth, arabinoxylan, or the like, or mixtures thereof.

In producing the liquid compositions of the present disclosure, there are preferred amounts of fiber, chondroitin sulfate, glucosamine, and calcium, when used. More particularly, it is preferred to use fiber in an amount ranging from greater than 0% to about 10% (w/w); chondroitin sulfate in an amount ranging from greater than 0% to about 0.5% (w/w); glucosamine in an amount ranger from greater than 0% to about 0.7% (w/w); and calcium in an amount ranging from greater than 0% to about 0.5% (w/w).

Any source of calcium is suitable for use in the liquid compositions of the disclosure. Examples of suitable sources of calcium include, but are not limited to, calcium lactate, calcium gluconate, a combination of calcium lactate and calcium gluconate, tricalcium phosphate, dicalcium phosphate, and the like, and mixtures thereof.

Suitable for use as buffers in the liquid compositions of the present disclosure, but not limited thereto, are potassium citrate, sodium citrate, potassium phosphate, sodium phosphate, and the like, and mixtures thereof.

5

10

15

20

25

30

Any sweetener may be used in the liquid compositions of the present disclosure. Examples of suitable sweeteners include, but are not limited to, high intensity sweeteners such as sucralose, acesulfame potassium, aspartame, stevia, thaumatin, neotame, MONATIN brand high intensity sweetener and the like, or mixtures thereof, corn syrups, high fructose corn syrups, sugar alcohols such as maltitol, erythritol, lactitol, xylitol, sucrose, fructose, dextrose, and the like, and mixtures thereof.

Any acid may be used in the liquid compositions of the present disclosure. Included in the suitable acids, but not limited thereto, are phosphoric acid, malic acid, citric acid, lactic acid and the like, and mixtures thereof.

Any fruit juice or fruit juice concentrate may be used in the liquid compositions of the disclosure. Exemplary fruit juice and fruit juice concentrate include, but are limited to, apple, orange, banana, strawberry, cranberry, kiwi, blueberry, raspberry, grape, mango, grapefruit, or the like. The fruit juices or fruit juice concentrate may be used alone or in combination.

The buffer, sweetener, fruit juice, fruit juice concentrate, and acid are used in any amount conventionally utilized in producing health-related liquid compositions of the type previously available.

The liquid compositions of the present disclosure may be prepared by any conventional technique. In the present disclosure, the liquid compositions of the Examples were prepared by the following process:

- (a) Water was metered, or weighed, into a mixing tank;
- (b) The fiber, which was inulin or beta-glucan in the Examples, was added to the water which was being agitated;
- (c) Thereafter, chondroitin sulfate, or glucosamine, or both, if used, was added to the water-inulin mixture;

- 3 -

(d) Thereafter the fruit juice or fruit juice concentrate, or both, was added to the mixture;

- (e) Then calcium lactate was added to the resultant mixture;
- (f) Then sodium citrate was added to the mixture;

5

10

15

20

25

30

- (g) The monopotassium phosphate was added to the mixture;
- (h) The sweeteners, which in the Examples included erythritol, trehalose, and acesulfame potassium, were added to the mixture;
- (i) Then flavors, which in the Examples were lemon, lime, or peach, were added to the mixture;
- (j) Then citric acid and malic acid were added to the mixture;
- (k) The pH of the mixture was measured and adjusted, if necessary, to a target pH of 4.0 ± 0.1 ; and
- (l) Then the mixture was filtered to provide the liquid composition of the disclosure.

The following examples are presented to illustrate the present disclosure and to assist one of ordinary skill in making and using the same. The examples are not intended in any way to otherwise limit the scope of the disclosure.

EXAMPLE 1

In this Example there is produced a lemon lime flavored health-related liquid composition, using the procedure set forth herein.

The liquid composition was prepared by mixing 94.7% (3585.0 g) water; 2.0% (75.7 g) erythritol; 1.4% (53 g) inulin, 0.78% (29.5 g) calcium lactate; 0.50% (18.9 g) trehalose; 0.08% (2.9 g) natural lemon flavor; 0.08% (2.9 g) natural lime flavor; 0.05% (1.9 g) citric acid; 0.05%(1.9 g) malic acid; 0.04% (1.5 g) sodium citrate, dihydrate; 0.02% (0.76 g) acesulfame potassium; 0.01% (0.4 g) monopotassium phosphate; and 0.17% (6.4 g) chondroitin sulfate. All percentages are weight/weight (w/w).

EXAMPLE 2

In this Example, there is produced a peach flavored health-flavored liquid composition, using the procedure set forth herein:

The liquid composition was prepared by mixing 94.7% (3585.8 g) water; 2.0% (75.7 g) erythritol; 1.4% (53 g) inulin, 0.78% (29.5 g) calcium lactate; 0.50% (18.9 g) trehalose; 0.13% (5 g) peach flavor; 0.05% (1.9 g) citric acid; 0.05% (1.9 g) malic acid;

0.04% (1.5 g) sodium citrate, dihydrate; 0.02% (0.76 g) acesulfame potassium; 0.01% (0.4 g) monopotassium phosphate; and 0.17% (6.4 g) chondroitin sulfate. All percentages are weight/weight (w/w).

EXAMPLE 3

In this Example, there is produced a health-related liquid composition comprising water, inulin, and chondroitin sulfate, using the procedure set forth herein.

5

10

15

20

25

30

The liquid composition was prepared by mixing 98.4% (4921.5 g) water; 1.4% (70 g) inulin; and 0.17% (8.5 g) chondroitin sulfate. All percentages are weight/weight (w/w)

EXAMPLE 4

In this Example, there is produced a health-related liquid composition comprising water, inulin, and glucosamine, using the procedure set forth herein.

The liquid composition was prepared by mixing 98.4% (4920.7 g) water; 1.4% (70 g) inulin; and 0.19% (9.4 g) glucosamine. All percentages are weight/weight (w/w)

EXAMPLE 5

In this Example, there is produced a health-related liquid composition comprising water, inulin, chondroitin sulfate, and glucosamine, using the procedure set forth herein.

The liquid composition was prepared by mixing 98.2% (4912.2 g) water; 1.4% (70 g) inulin; 0.17% (8.5 g) chondroitin sulfate; and 0.19% (9.4 g) glucosamine. All percentages are weight/weight (w/w)

EXAMPLE 6

In this Example, there is produced a health-related liquid composition comprising water, inulin, chondroitin sulfate, glucosamine, and calcium lactate, using the procedure set forth herein.

The liquid composition was prepared by mixing 97.5% (4873.2 g) water; 1.4% (70 g) inulin; 0.17% (8.5 g) chondroitin sulfate; 0.19% (9.4 g) glucosamine; and 0.78% (39 g) calcium lactate. All percentages are weight/weight (w/w)

EXAMPLE 7

In this Example, there is produced a health-related fruit juice-containing liquid composition comprising water, cranberry juice concentrate, white grape juice concentrate, inulin, chondroitin sulfate, glucosamine, and calcium lactate, using the procedure set forth herein.

The liquid composition was prepared by mixing 76.2% (3811.7 g) water; 4.1% (202.5 g) cranberry juice concentrate; 17.2% (859 g) white grape juice concentrate; 1.4% (70 g) inulin; 0.17% (8.5 g) chondroitin sulfate; 0.19% (9.4 g) glucosamine; and 0.78% (39 g) calcium lactate. All percentages are weight/weight (w/w)

5

10

EXAMPLE 8

In this Example, there is produced a health-related fruit juice-containing liquid composition comprising orange juice, beta-glucan fiber, chondroitin sulfate, glucosamine, and calcium lactate, using the procedure set forth herein.

The liquid composition was prepared by mixing 98.3% (4916.4 g) orange juice, that includes water, 0.54% (26.7 g) beta-glucan, 0.17% (8.5 g) chondroitin sulfate; 0.19% (9.4 g) glucosamine; and 0.78% (39 g) calcium lactate. All percentages are weight/weight (w/w).

All of the liquid compositions of Examples 1 through 8 were evaluated for taste, and were found to be acceptable for consumption.

This disclosure has been described with reference to various specific and illustrative embodiments and techniques. However, one skilled in the art will recognize that many variations and modifications may be made while retaining within the spirit and scope of the disclosure and the claims.

CLAIMS

We claim:

15

25

1. A liquid composition comprising water, a fiber, and a glycosaminoglycan selected from the group consisting of chondroitin sulfate, glucosamine, and mixtures thereof.

- 5 2. The liquid composition according to Claim 1 wherein the fiber is present in an amount of greater than zero (0) to about 10% (weight/weight), the chondroitin sulfate is present in an amount of greater than zero (0) to about 0.5% weight/weight, and the glucosamine is present in an amount of greater than zero (0) to about 0.7% weight/weight.
- 3. The liquid composition according to Claim 1 wherein the fiber is selected from the group consisting of inulin, polydextrose, arabinogalactose, hydrolyzed guar gum, gum arabic, locust bean gum, xanthan, pectin, alginate, carageenan, beta-glucans, tragacanth, arabinoxylan, and mixtures thereof.
 - 4. The liquid composition according to Claim 1 wherein the fiber is inulin.
 - 5. The liquid composition according to Claim 1 wherein the glycosaminoglycan is chondroitin sulfate.
 - 6. The liquid composition according to Claim 1 wherein the fiber is inulin and the glycosaminoglycan is chondroitin sulfate.
 - 7. The liquid composition according to Claim 1 wherein the glycosaminoglycan is glucosamine.
- 20 8. The liquid composition according to Claim 1 wherein the fiber is inulin and the glycosaminoglycan is glucosamine.
 - 9. The liquid composition according to Claim 1 wherein the glycosaminoglycan is a mixture of chondroitin sulfate and glucosamine.
 - 10. The liquid composition according to Claim 1 wherein the fiber is inulin, and the glycosaminoglycan is a mixture of chondroitin sulfate and glucosamine.
 - 11. The liquid composition according to Claim 1 further comprising a source of calcium.
 - 12. The liquid composition according to Claim 11 wherein the source of calcium is present in an amount of greater than zero (0) to about 0.5% weight/weight.
- 30 13. The liquid composition according to Claim 6 further comprising a source of calcium.

14. The liquid composition according to Claim 8 further comprising a source of calcium.

- 15. The liquid composition according to Claim 10 further comprising a source of calcium.
- 5 16. The liquid composition according to Claim 1 further comprising a component selected from the group consisting of at least one sweetener, a source of calcium, at least one flavor additive, at least one acid, sodium citrate, trehalose, monopotassium phosphate, at least one fruit juice, at least one fruit juice concentrate, and mixtures thereof.
- 17. The liquid composition according to Claim 16 wherein the sweetener comprises a mixture of erythritol and acesulfame potassium, the source of calcium comprises calcium lactate, and the acid comprises a mixture of citric acid and malic acid.
 - 18. The liquid composition according to Claim 17 wherein the fiber is inulin.
 - 19. The liquid composition according to Claim 18 wherein the glycosaminoglycan is chondroitin sulfate.
- 15 20. The liquid composition according to Claim 18 wherein the glycosaminoglycan is glucosamine.
 - 21. The liquid composition according to Claim 18 wherein the glycosaminoglycan is a mixture of chondroitin sulfate and glucosamine.
- The liquid composition according to Claim 16 wherein the sweetener comprises amixture of erythritol and high fructose corn syrup.

INTERNATIONAL SEARCH REPORT

International application No PCT/US2006/007921

A. CLASSII INV.	FICATION OF SUBJECT MATTER A61K31/726 A61K31/7008 A23L2/52	A23L1/05						
	hinternational Patent Classification (IPC) or to both national classifica	tion and IPC						
	SEARCHED cumentation searched (classification system followed by classification	on symbols)						
A61K	A23L	•						
Documentat	ion searched other than minimum documentation to the extent that so	uch documents are included in the fields se	arched					
Electronic d	ata base consulted during the international search (name of data bas	e and, where practical, search terms used)					
EPO-Internal, WPI Data								
C. DOCUME	ENTS CONSIDERED TO BE RELEVANT							
Category*	Citation of document, with indication, where appropriate, of the rele	evant passages	Relevant to claim No.					
χ	ANONYMOUS: "Liquid		1,3,5,7,					
į	Glucosamine/Chondroitin/MSM - 16	oz"	9,16					
	INTERNET ARTICLE, 11 February 2005 (2005-02-11), pa	~~ 1 0						
	XP002392912	ges 1-2,						
	Retrieved from the Internet:							
(URL:http://web.archive.org/web/20							
	118/www.nowfoods.com/?action=item	detail⁢						
	em_id=2841> page 1							
χ	US 2003/069202 A1 (KERN KENNETH N	ORMAN ET	1-22					
	AL) 10 April 2003 (2003-04-10)							
	paragraphs [0002], [0007], [004 [0055], [0107], [0117], [0120]	/] -						
	[0121]; examples 7,18-20	,						
		,						
		/						
	<u> </u>							
	ner documents are listed in the continuation of Box C.	X See patent family annex.						
		"T" later document published after the inte or priority date and not in conflict with						
"A" docume consid	ent defining the general state of the art which is not ered to be of particular relevance	cited to understand the principle or the invention						
"E" earlier o	document but published on or after the international ate	"X" document of particular relevance; the c						
"L" docume	nt which may throw doubts on priority claim(s) or is cited to establish the publication date of another	involve an inventive step when the do	cument is taken alone					
citation	n or other special reason (as specified)	"Y" document of particular relevance; the c cannot be considered to involve an involve an involve and	entive step when the					
other r		document is combined with one or mo ments, such combination being obviou in the art.						
	ent published prior to the international filing date but an the priority date claimed	*& document member of the same patent	family					
Date of the	actual completion of the international search	Date of mailing of the international sea	rch report					
1	August 2006	22/08/2006						
Name and n	nailing address of the ISA/	Authorized officer						
	European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl,							
	Fax: (+31-70) 340-3016	Rinaldi, F						

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2006/007921

C(Continua	ation). DOCUMENTS CONSIDERED TO BE RELEVANT		
ategory*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	
X	US 2001/051134 A1 (PANDYA MAHENDRA) 13 December 2001 (2001-12-13) paragraphs [0011], [0013], [0024], [0035], [0044]; claims 8,24	1-22	
Х	WO 99/18934 A (VANDERBILT UNIVERSITY) 22 April 1999 (1999-04-22) example 3	1,3,5, 11,12,16	
A	example 3 DE 101 09 798 A1 (AVENTIS PHARMA DEUTSCHLAND GMBH) 12 September 2002 (2002-09-12) paragraph [0188]	1-22	

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/US2006/007921

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
US 2003069	202 A1	10-04-2003	NONE		
US 2001051	134 A1	13-12-2001	NONE		
WO 9918934	A	22-04-1999	AU EP	9799198 A 1021168 A1	03-05-1999 26-07-2000
DE 1010979	8 A1	12-09-2002	NONE		